K011286

MAY 16 2001

510(k) SUMMARY

Submitted For:

NINGBO YUJIANG PLASTIC & RUBBER CO., LTD
Economic & Tech Development Zone G-4
Ningbo, Zhejiang Province
315803, P.R. China

Submitted by:

TUCKER & ASSOCIATES
Official Correspondent and United States Agent for
Ningbo Yujiang Plastic & Rubber Co., Ltd.
JANNA P. TUCKER, President – CEO
198 Avenue de la D'emerald
Sparks, NV 89434
Phone: 775-342-2612 Fax: 775-342-2613
E-mail: Tuckerjan@aol.com

Date of Submission:

Device Name:

POWDER-FREE NITRILE EXAM GLOVE, BLUE
Class I Device, 80LZA

Proprietary Name:

(Multiple Labels) Powder-Free Nitrile Exam Gloves, Blue

Labels/Labeling:

This device will be marketed to healthcare professionals at Dentist and Doctor Offices, Laboratories, Clinics and Hospitals through its intended use.

Intended Use:

A patient examination glove is a disposable device intended For medical purposes that is worn on the examiner's hand Or finger to prevent contamination between patient and Examiner.

Substantial Equivalence:

This device is equivalent to those in commercial distribution. They are to be worn as a protective device on the examiner's hand or finger, also protecting the patient.

Both in its intended use and/or physical characteristics, this Device is equivalent to devices currently marketed by U.S. Companies. It is **substantially equivalent** to the devices Manufactured by Shanghai Poseidon (K001128) and Ningbo Shasta Asia Partners (K000670)

Test Results (Means

EXHIBIT N
Page 52 of 53

And/or Results):	This device has met or exceeded the following Standards/Tests: ASTM D 3578-00 ASTM D 5712-99 ASTM D 5151-00 ASTM D 6124-00 BIO-BURDEN ISO 2859 Bio-Compatibility: Dermal Sensitization Primary Skin Irritation
Conclusion:	This device is substantially equivalent to the devices Approved as K001128 and K000670.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

MAY 1 6 2001

Ningbo Yujiang Plastic & Rubber Company Limited
C/O Ms. Janna P. Tucker
Official Correspondent
Janna Tucker & Associates
198 Avenue De La D' Emerald
Sparks, Nevada 89434-9550

Re: K011286
Trade/Device Name: Powder-Free Nitrile Examination Glove, Blue
Regulation Number: 880.6250
Regulatory Class: I
Product Code: LZA
Dated: April 25, 2001
Received: April 27, 2001

Dear Ms. Tucker:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements

concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4692. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "http://www.fda.gov/cdrh/dsma/dsmamain.html".

Sincerely yours,

Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

INDICATIONS FOR USE

APPLICANT: **NINGBO YUJIANG PLASTIC & RUBBER CO., LTD.**

510(k) NUMBER: K011286

DEVICE NAME: **POWDER-FREE NITRILE EXAM GLOVES, BLUE**

A patient examination glove is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use____ OR Over-The-Counter Use____
(Per 21 CFR 801.109)

(Optional Format 1-2-96)

(Division Sign-Off)
Division of Dental, Infection Control, General Hospital Devices
510(k) Number K011286

EXHIBIT B
Page 2 of 53